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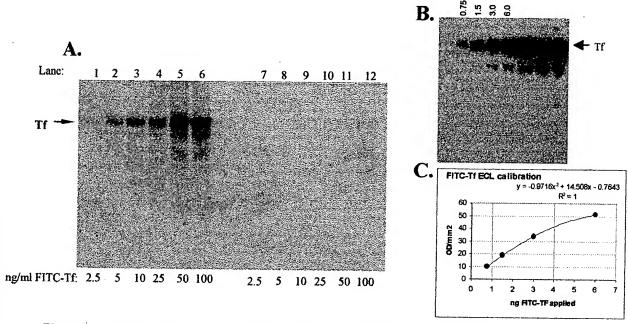
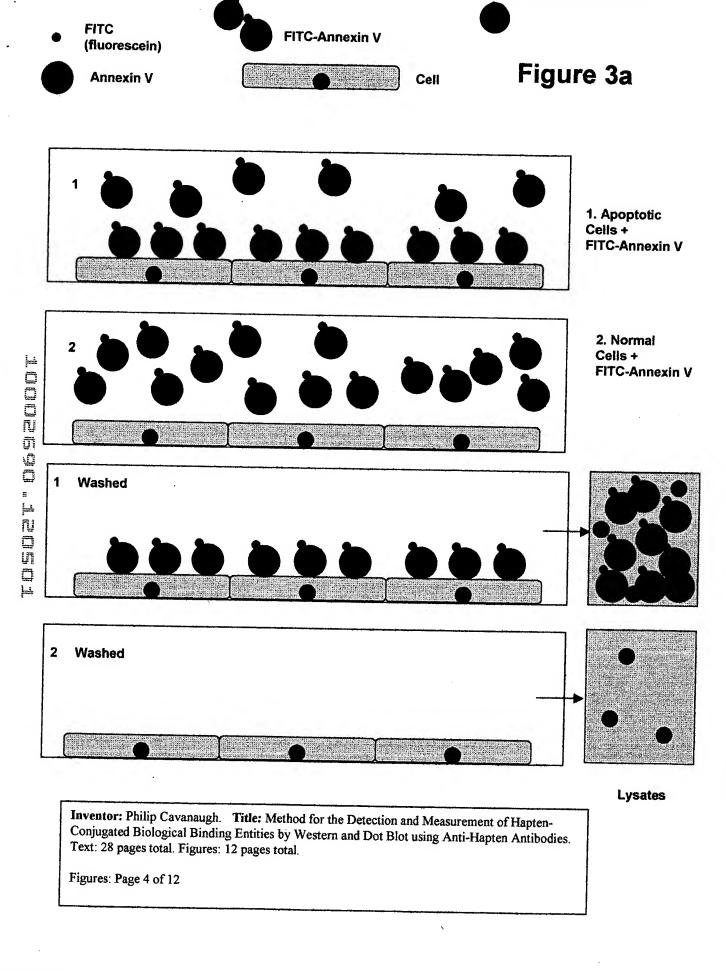
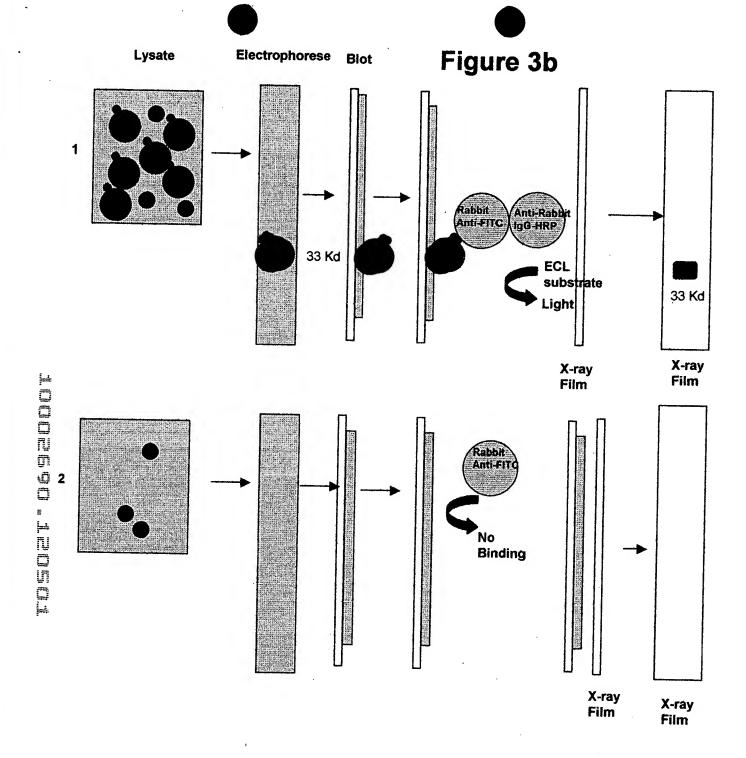


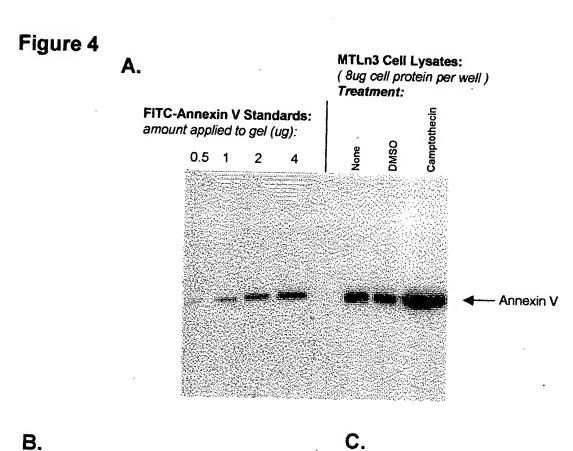
Figure 2. A. Measurement of FITC-transferrin (FITC-Tf) binding to MTLn2/TfR cells. Cells at 50 -60 % confluency growing in 12 well plates were serum-starved, then treated at 4 C with increasing levels of FITC-Tf. After 2h, cells were washed, lysed, and equal cell equivalents were electrophoresed, blotted, incubated with goat anti-FITC, then with anti-goat-HRP and an HRP ECL substrate. The blot was then analyzed using the BIO-RAD fluor S Multimager. Lanes 1-6 = lysates from cell exposed to the concentration of FITC-TF listed below the blot. Lanes 7-12 = lysates from cells treated as in 1-6, but also with a 200-fold excess of unconjugated Tf. B. Results from a blot treated as in A but loaded with pure FITC-TF standards in the amounts (in ng) indicated on the top. C. Analysis of the bands from B, indicating the type of standard curves achievable.

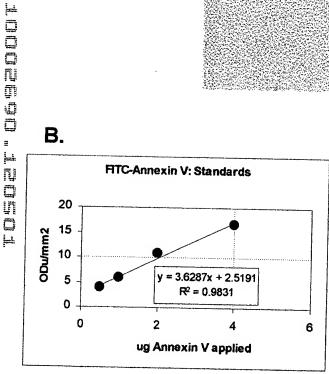
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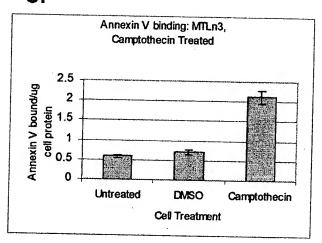




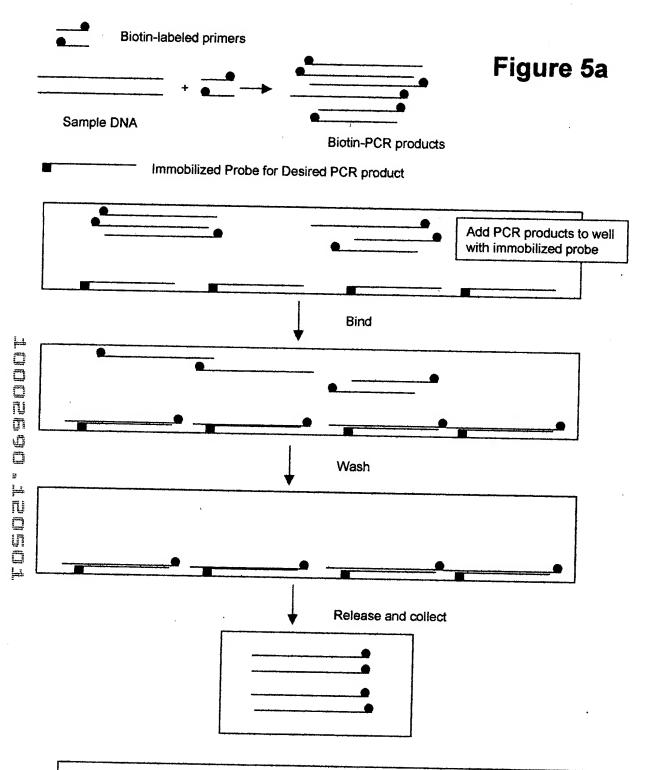
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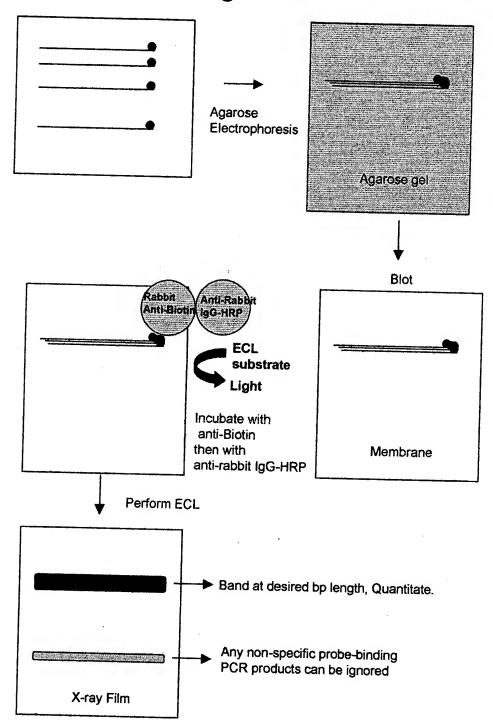


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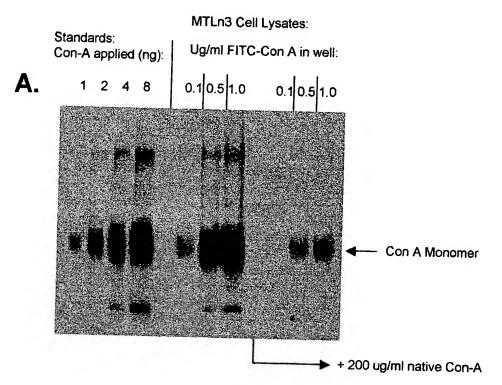
Figure 5b

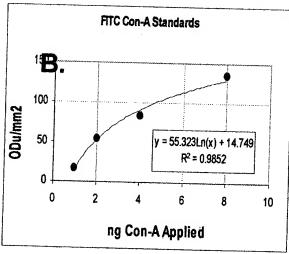


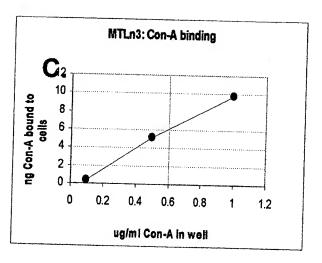
Inventor: Philip Cavanaugh. Title: Method for the Detection and Measurement of Hapten-Conjugated Biological Binding Entities by Western and Dot Blot using Anti-Hapten Antibodies. Text: 28 pages total. Figures: 12 pages total.

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Figure 6.



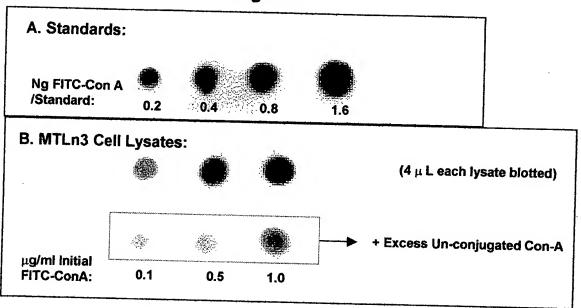


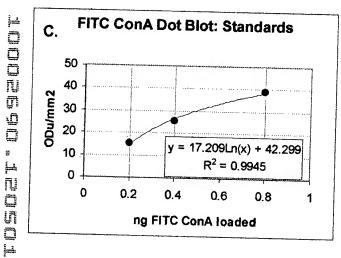


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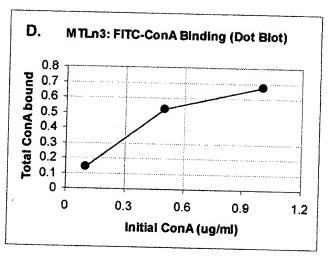
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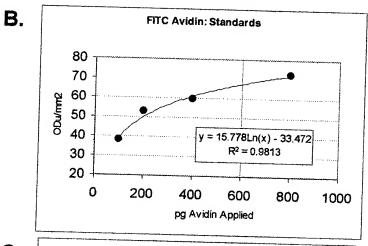
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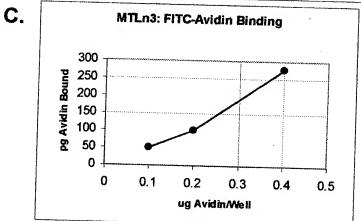
Figure 8

A. FITC-Avidin Standards, pg loaded onto gel:
100 200 400 800

0.1 0.2 0.4

Avidin





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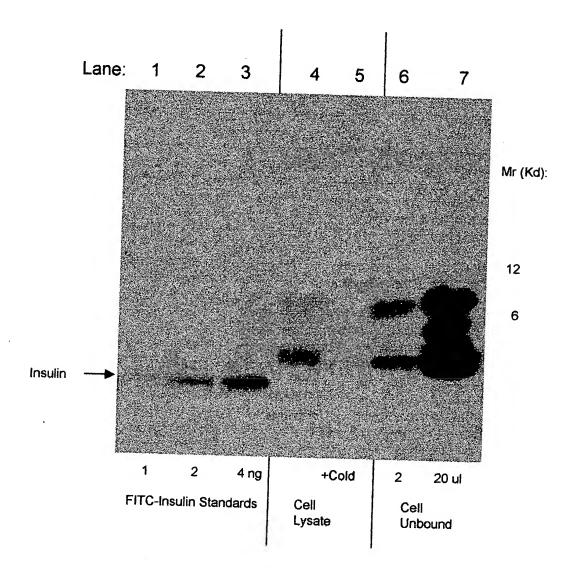


Figure 9: FITC-Insulin binding by K562 cells.

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